

Research Article

Complete mitogenome of the critically endangered Asian king vulture (*Sarcogyps calvus*) (Aves, Accipitriformes, Accipitridae): evolutionary insights and comparative analysis

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Abstract

The Asian king vulture (Sarcogyps calvus), also known as the red-headed vulture, is an Old World vulture (Gypini) facing severe population declines. This study aimed to assemble the complete mitogenome of S. calvus, explore its phylogenetic relationships, estimate divergence times, and examine genetic distances and amino acid substitutions. The mitogenome was de novo assembled from genomic DNA extracted from the blood of a female S. calvus. Phylogenetic and pairwise genetic distance analyses were conducted, comparing S. calvus with other members of Gypini, New World vultures (Cathartidae) and various other birds. The assembled mitogenome was 17,750 base pairs in length, comprising 13 protein-coding genes (PCGs), 22 transfer RNA genes, two ribosomal RNA genes and two control regions. Most PCGs used the ATG start codon, except for cytochrome c oxidase subunit 1 (COX1), which employed GTG. Phylogenetic analysis revealed a close genetic relationship between S. calvus and other members of Gypini, with an estimated divergence time of 16.7 million years ago. Genetic distance analysis indicated that S. calvus was more closely related to other Gypini, as well as to Spilornis cheela and Circaetus pectoralis (Circaetini)), than to Cathartidae. Conserved amino acid substitutions between Gypini and Cathartidae were primarily observed in the NADH-ubiquinone oxidoreductase chain 1 (ND1) gene. This study provided the first complete mitogenome of S. calvus, offering new insights into its genomic structure, evolutionary history, and genetic relationships.

Key words: Asian king vulture, comparative analysis, evolution, mitogenome, *Sarcogyps* calvus

Introduction

The Asian king vulture (*Sarcogyps calvus* Scopoli, 1786), also known as the red-headed vulture, belongs to the Old World vulture group (tribe Gypini) within the order Accipitriformes and the family Accipitridae. Classified as Critically Endangered by the International Union for Conservation of Nature (IUCN) Red List of Threatened Species, *S. calvus* is also listed under Appendix II of the Convention on International Trade in Endangered Species of Wild Fauna and Flora (CITES) (BirdLife International 2021; CITES 2024). Additionally, the species is protected under Thailand's Wild Animal Conservation and Protection Act, B.E. 2562 (2019) (FAOLEX Database 2019). Gypini are distributed across Africa, Asia, and Europe, while New World vultures (Cathartidae) inhabit the Americas. Currently, the global population of *S. calvus* in the wild is estimated to be between 2,500 and 9,999 individuals, while in Thailand, the species is possibly extinct in the wild and only seven individuals remain in captivity (BirdLife International 2021; Buthasane et al. 2024).

The mitogenome (mitochondrial genome) is a valuable tool for investigating phylogenetic relationships, molecular identification, and adaptive evolution (De Panis et al. 2021; Kong et al. 2024). Mitogenomes of four species of Gypini have been reported, i.e., those of the Eurasian griffon (Gyps fulvus Hablizl, 1783), cape vulture (Gyps coprotheres Forster, 1798), cinereous vulture (Aegypius monachus Linnaeus, 1766) and Himalayan griffon (Gyps himalayensis Hume, 1869) (Li et al. 2015; Mereu et al. 2017; Jiang et al. 2019; Adawaren et al. 2020). For Cathartidae, seven mitogenomes from six species have been documented, i.e. the Andean condor (Vultur gryphus Linnaeus, 1758), king vulture (Sarcoramphus papa Linnaeus, 1758), California condor (Gymnogyps californianus Shaw, 1797), lesser yellow-headed vulture (Cathartes burrovianus Cassin, 1845), turkey vulture (Cathartes aura Linnaeus, 1758) and black vulture (Coragyps atratus Bechstein, 1793) (Slack et al. 2007; De Panis et al. 2021; Urantówka et al. 2021). In addition, 11 mitogenomes from other species in the family Accipitridae have been reported, including those of the golden eagle (Aquila chrysaetos Linnaeus, 1758), common buzzard (Buteo buteo Linnaeus, 1758) and black kite (Milvus migrans Boddaert, 1783) (Haring et al. 2001; Jeon et al. 2018; Mead et al. 2021). The genome of S. calvus has recently been published (Buthasane et al. 2024). However, no mitogenomic data are currently available for S. calvus, and its mitochondrial features remain understudied. This study aimed to elucidate the complete mitogenome of S. calvus and provide a comprehensive analysis of its structure, phylogenetic position, and the divergence time from other vultures. This research offers valuable insights into the mitochondrial profiles, evolutionary relationships, and population genetics of S. calvus in relation to other Gypini, Cathartidae and related species.

Materials and methods

Whole blood samples were obtained from a female *S. calvus*, approximately 25 years old, at Nakhon Ratchasima Zoo, the Zoological Park Organization of Thailand (ZPOT). Sampling was conducted in compliance with the ethical guidelines under the Chulalongkorn University Animal Care and Use Committee

(CU-ACUC), Thailand (approval number 2131005). Total DNA was extracted from the whole blood sample using the Wizard HMW DNA Extraction Kit (Promega, Madison, WI, USA). The DNA concentration was determined using a NanoDrop One Microvolume UV-Vis Spectrophotometer (Thermo Fisher Scientific, Waltham, MA, USA).

The mitochondrial genome was sequenced using the short-read MGIS-EQ-2000 platform (MGI Tech, Shenzhen, China) and assembled with NOVOPlasty v. 3.8.2 (Dierckxsens et al. 2017). Annotation was carried out using the MITOS WebServer (Bernt et al. 2013). Protein-coding, rRNA and tRNA genes were further identified using the NCBI Basic Local Alignment Search Tool (BLAST) (Altschul et al. 1990). The circular structure of the mitogenome was visualized with OrganellarGenomeDRAW (OGDRAW) v. 1.3.122 (Greiner et al. 2019). Analyses of nucleotide and amino acid composition were conducted using MEGA X (Kumar et al. 2018). Simple sequence repeats (SSRs) of 1–6 bp in length were identified using the Microsatellite identification tool (Beier et al. 2017).

Mitogenomes from 39 bird species, representing the orders Accipitriformes (Old World vultures, hawks, eagles, and kites), Cathartiformes (New World vultures), Falconiformes (falcons), Strigiformes (owls), Anseriformes (ducks and relatives) and Galliformes (chickens and relatives), were used for phylogeny reconstruction, comparative codon usage analyses (Table 1) and divergence time estimation. These mitogenomes, along with the newly sequenced mitogenome of S. calvus, were analyzed. Multiple sequence alignments of 13 conserved protein-coding genes (PCGs)-cytochrome B (CYTB), NADH dehydrogenase subunits 1 (ND1), ND2, ND3, ND4, ND4L, ND5, ND6, cytochrome c oxidase subunits 1 (COX1), COX2, COX3, ATP synthase F0 subunit 6 (ATP6) and ATP8-were performed using PRANK v170427. The best-fit model, mt-Ver+I+R4, was selected using ModelFinder, and maximum likelihood phylogenies were constructed using IQ-TREE v. 2.2.0.3 with 1,000 ultrafast bootstrap replications (Nguyen LT et al. 2014; Hoang et al. 2017). The phylogenetic tree was visualized with Figtree v. 1.4.4 (Rambaut 2018). Several species of Anseriformes and Galliformes were used as outgroups. Divergence times were estimated using MCMCTree in the PAML 4.9j package, with the Hessian matrix computed via CODEML and a burn-in of 2,000 iterations. Fossil calibration times were obtained from the TimeTree database (Kumar et al. 2022). Genetic distance analyses were conducted using MEGA X and visualized using ggplot2 and ggtree in R (Wickham 2016; Yu et al. 2017; Kumar et al. 2018). The nomenclature for higher taxa follows Gregory et al. (2024).

Relative synonymous codon usage (RSCU) values for *S. calvus* mitochondrial protein-coding genes were calculated using MEGA X (Kumar et al. 2018). RSCU values reflect codon bias, with values greater than one indicating positive codon bias, values less than one indicating a negative codon bias, and values equal to one indicating random codon usage (Wong et al. 2010).

Amino acid sequences from 13 mitochondrial protein-coding genes in the mitogenomes of Gypini (Aegypius monachus, Gyps coprotheres, Gyps fulvus, Gyps himalayensis, S. calvus) and Cathartidae (Cathartes aura, Cathartes burrovianus, Coragyps atratus, Gymnogyps californianus, Sarcoramphus papa, Vultur gryphus) were aligned using the Unipro UGENE Multiple Alignment Editor (Okonechnikov et al. 2012).

Table 1. List of 39 avian species employed for comparative mitogenome analyses in this study, along with their corresponding GenBank accession numbers.

Scientific name	Order	Family	Accesion number	Sequence length (bp)	Reference
Accipiter gentilis	Accipitriformes	Accipitridae	NC_011818.1	18,266	Unpublished
Accipiter virgatus	Accipitriformes	Accipitridae	NC_026082.1	17,952	Song et al. 2015
Aegypius monachus	Accipitriformes	Accipitridae	KF682364.1	17,811	Li et al. 2015
Aquila chrysaetos	Accipitriformes	Accipitridae	NC_024087.1	17,332	Doyle et al. 2014
Buteo buteo	Accipitriformes	Accipitridae	NC_003128.3	18,674	Haring et al. 2001
Circaetus pectoralis	Accipitriformes	Accipitridae	NC_052805.1	17,473	Feng et al. 2020
Circus melanoleucos	Accipitriformes	Accipitridae	NC_035801.1	17,749	Unpublished
Gyps coprotheres	Accipitriformes	Accipitridae	MF683387.1	16,908	Adawaren et al. 2020
Gyps fulvus	Accipitriformes	Accipitridae	NC_036050.1	18,094	Mereu et al. 2017
Gyps himalayensis	Accipitriformes	Accipitridae	KY594709.1	17,381	Jiang et al. 2019
Haliaeetus albicilla	Accipitriformes	Accipitridae	NC_040858.1	17,719	Kim et al. 2019
Haliastur indus	Accipitriformes	Accipitridae	NC_066800.1	19,055	Sonongbua et al. 2024
Milvus migrans	Accipitriformes	Accipitridae	NC_038195.1	18,016	Jeon et al. 2018
Spilornis cheela	Accipitriformes	Accipitridae	NC_015887.1	18,291	Unpublished
Spizaetus tyrannus	Accipitriformes	Accipitridae	NC_052803.1	17,479	Feng et al. 2020
Pandion haliaetus	Accipitriformes	Pandionidae	NC_008550.1	19,285	Gibb et al. 2007
Sagittarius serpentarius	Accipitriformes	Sagittariidae	NC_023788.1	19,329	Mahmood et al. 2014
Anser cygnoides	Anseriformes	Anatidae	NC_023832.1	19,302	Mu et al. 2014
Branta canadensis	Anseriformes	Anatidae	NC_007011.1	16,808	Snyder et al. 2015
Anseranas semipalmata	Anseriformes	Anseranatidae	NC_005933.1	16,870	Harrison et al. 2004
Cathartes aura	Cathartiformes	Cathartidae	NC_007628.1	16,870	Slack et al. 2007
Cathartes burrovianus	Cathartiformes	Cathartidae	NC_063526.1	16,779	Urantówka et al. 2021
Coragyps atratus	Cathartiformes	Cathartidae	NC_063525.1	17,864	Urantówka et al. 2021
Gymnogyps californianus	Cathartiformes	Cathartidae	BK059163.1	16,760	De Panis et al. 2021
Sarcoramphus papa	Cathartiformes	Cathartidae	NC_063527.1	16,773	Urantówka et al. 2021
Vultur gryphus	Cathartiformes	Cathartidae	NC_058600.1	16,739	De Panis et al. 2021
Caracara plancus	Falconiformes	Falconidae	NC_044672.1	17,077	Oswald et al. 2019
Falco peregrinus	Falconiformes	Falconidae	NC_000878.1	18,068	Mindell et al. 1997
Alectura lathami	Galliformes	Megapodiidae	NC_007227.1	16,698	Slack et al. 2007
Crax rubra	Galliformes	Cracidae	NC_024618.1	16,666	Meiklejohn et al. 2014
Callipepla squamata	Galliformes	Odontophoridae	NC_029340.1	16,701	Halley et al. 2015
Gallus gallus	Galliformes	Phasianidae	NC_053523.1	16,784	Unpublished
Numida meleagris	Galliformes	Numididae	NC_034374.1	16,785	Unpublished
Asio otus	Strigiformes	Strigidae	NC_039736.1	17,555	Lee et al. 2018
Bubo bubo	Strigiformes	Strigidae	NC_038219.1	18,952	Kang et al. 2018
Glaucidium cuculoides	Strigiformes	Strigidae	NC_034296.1	17,392	Unpublished
Otus sunia	Strigiformes	Strigidae	NC_041422.1	17,835	Zhou et al. 2019
Strix uralensis	Strigiformes	Strigidae	NC_038218.1	18,708	Kang et al. 2018
Phodilus badius	Strigiformes	Tytonidae	NC_023787.1	17,086	Mahmood et al. 2014

Results

The complete mitogenome of *S. calvus* was determined to be 17,750 base pairs (bp) in length and was assigned GenBank accession number OR896160. The circular structure of the mitogenome of *S. calvus* is illustrated in Fig. 1. This mitogenome contained 13 PCGs, 22 transfer RNA genes (tRNAs), two ribosomal RNA genes and two putative control regions (CRs), also referred to as D-loop regions (Table 2). The nucleotide composition was characterized by 54.1% adenine and thymine (AT) and 45.9% guanine and cytosine (GC).

The protein-coding regions spanned 11,407 bp, accounting for 64.26% of the length of the complete mitogenome of *S. calvus*. All PCGs, except for *ND6*, were transcribed on the plus strand. The predominant start codon for most PCGs

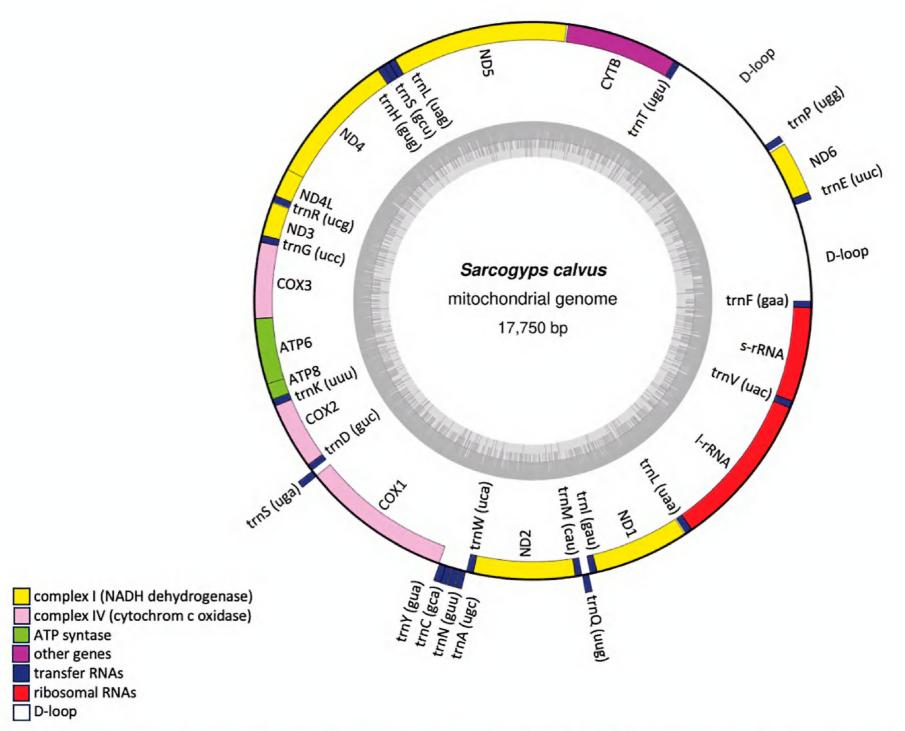


Figure 1. Circular mitogenome map of Asian king vulture. The complex I (NADH dehydrogenase), complex IV (cyto-chrome c oxidase), ATP synthase, ribosomal RNAs, transfer RNAs, cytochrome b and control region (D-loop) are annotated. Genes located outside the circle are transcribed in a clockwise direction, whilefig. genes inside are transcribed counterclockwise. The inner ring shadow denotes the GC content of the genome.

was ATG, except for *COX1*, which utilized GTG as the start codon (Table 2). A detailed overview of the RSCU and codon distribution in the protein-coding genes of the mitogenome of *S. calvus* is provided. The codons CUA (L), CCU (P), and CUC (L) exhibited the highest frequency of occurrence (Fig. 2). A total of 1,257 SSRs were identified in the mitogenome, comprising 314 (24.98%) mono-, 529 (42.08%) di-, 301 (23.95%) tri-, 78 (6.21%) tetra-, 24 (1.91%) penta- and 11 (0.88%) hexanucleotide repeats. The ND5 gene contained the highest number of SSRs with 135 repeats (Table 3).

The mitogenome of *S. calvus* was aligned with 39 previously published mitogenomes of bird species from the orders Accipitriformes, Cathartiformes, Falconiformes, Strigiformes, Anseriformes and Galliformes. Maximum likelihood phylogenies are illustrated in Fig. 3. The mitogenome of *S. calvus* was part of a clade formed by the tribe Gypini (*Gyps fulvus*, *Gyps coprotheres*, *Gyps himalayensis* and *Aegypius monachus*). Gypini formed a sister group with the serpent-eagles of the tribe Circaetini (*Spilornis cheela* Latham, 1790 and *Circaetus pectoralis* Smith, 1829). Gypini and Circaetini formed the sister-group of a clade comprising the subfamilies Accipitrinae and Aquilinae. The subfamily Aquilinae included the species *Spizaetus tyrannus* Wied, 1820 and *Aquila chrysaetos*, whereas the subfamily Accipitrinae included the tribe Accipitrini

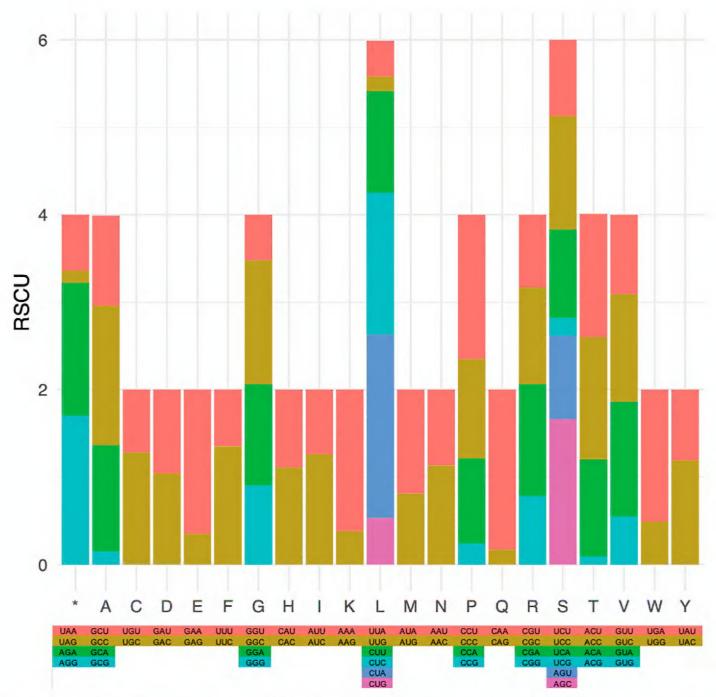


Figure 2. The relative synonymous codon usage (RSCU) and codon distribution of the Asian king vulture mitogenome. The different colors in the column chart symbolize distinct codon families associated with the amino acids listed below. Consistent coloring is applied to maintain representation uniformity across the same codon families. Bar chart showed-relative synonymous codon usage in all protein-coding genes of the mitogenome of *S. calvus*.

(Accipiter virgatus Temminck, 1822, Accipiter gentilis and Circus melanoleucos Forster, 1795) and the tribe Buteonini (Milvus migrans, Haliastur indus Boddaert, 1783, Haliaeetus albicilla and Buteo buteo) (Fig. 3A). The estimated evolutionary divergence time between S. calvus and its sister taxa, based on mitochondrial data, was approximately 22.2 million years ago (Mya) (95% highest posterior density (HPD): 2.8–43.8 Mya) (Fig. 3B). Fossil calibration constraints were applied to several groups, including Accipitriformes and Falconiformes, as well as Accipitriformes and Galliformes, among others. The estimated divergence times for these groups ranged from 6.2 to 101.5 Mya. Genetic distance analysis showed that the genetic distance between S. calvus and other Gypini ranged from 4.02% to 5.17%, while the distance between S. calvus and Cathartidae ranged from 10.90% to 12.26% (Suppl. material 1: table S1).

A total of 138 conserved amino acid substitution sites were observed between Old World vulture (Gypini) and New World vulture (Cathartidae) groups. The largest number of substitutions was found in the *ND5* gene (23 sites), whereas the smallest number occurred in *ATP6* (3 sites) (Suppl. material 1: table S2). Among these, 67 sites displayed substitutions between different amino acid chemical groups, with *ND1* showing the highest number of such substitutions (11 sites). Special case amino acid substitutions were found in *ATP8* (1 site), *COX1* (1 site), *COX2* (1 site), *CYTB* (2 sites), *ND1* (2 sites), *ND3* (2 sites),

 Table 2. Characteristics of the mitogenome of Sarcogyps calvus.

Start	End	Length (bp)	Direction	Туре	Gene name	Gene product	Anti-codon	Start codon	Stop codon
1	987	987	+	control region	-	-	_	_	_
988	1058	71	+	tRNA	trnE(uuc)	tRNA-Glu	TTC	_	_
1059	1577	519	+	CDS	ND6	NADH dehydrogenase subunit 6	_	ATG	TAG
1599	1668	70	+	tRNA	trnP(ugg)	tRNA-Pro	TGG	-	-
1669	2863	1195	+	control region					
2931	2864	68	_	tRNA	trnT(ugu)	tRNA-Thr	TGT		
4076	2934	1143	-	CDS	СҮТВ	cytochrome b	_	ATG	TAA
5903	4089	1815	_	CDS	ND5	NADH dehydrogenase subunit 5	-	ATG	TAA
5974	5904	71	_	tRNA	trnL(uag)	tRNA-Leu	TAG	_	_
6039	5975	65	_	tRNA	trnS(gcu)	tRNA-Ser	GCT	_	_
6110	6041	70	_	tRNA	trnH(gug)	tRNA-His	GTG	_	_
7488	6106	1383	_	CDS	ND4	NADH dehydrogenase subunit 4	_	ATG	AGG
7778	7482	297	_	CDS	ND4L	NADH dehydrogenase subunit 4L	_	ATG	TAA
7848	7780	69	_	tRNA	trnR(ucg)	tRNA-Arg	TCG	_	_
8204	7854	354	_	CDS	ND3	NADH dehydrogenase subunit 3	_	ATG	AGG
8273	8205	69	_	tRNA	trnG(ucc)	tRNA-Gly	TCC	-	-
9057	8274	784	_	CDS	COX3	cytochrome c oxidase subunit III	-	ATG	CCT
9740	9057	684	_	CDS	ATP6	ATP synthase F0 subunit 6	_	ATG	TAA
9898	9731	168	<u> </u>	CDS	ATP8	ATP synthase F0 subunit 8	_	ATG	TAA
9970	9900	71	-	tRNA	trnK(uuu)	tRNA-Lys	TTT	-	_
10655	9972	684	_	CDS	COX2	cytochrome c oxidase subunit II	_	ATG	TAA
10726	10658	69	_	tRNA	trnD(guc)	tRNA-Asp	GTC	-	_
10731	10802	72	+	tRNA	trnS(uga)	tRNA-Ser	TGA	_	-
12344	10794	1551	_	CDS	COX1	cytochrome c oxidase subunit I	_	GTG	AGG
12346	12415	70	+	tRNA	trnY(gua)	tRNA-Tyr	GTA	_	-
12416	12482	67	+	tRNA	trnC(gca)	tRNA-Cys	GCA	_	_
12485	12557	73	+	tRNA	trnN(guu)	tRNA-Asn	GTT	_	-
12560	12628	69	+	tRNA	trnA(ugc)	tRNA-Ala	TGC	-	-
12702	12630	73	-	tRNA	trnW(uca)	tRNA-Trp	TCA	_	-
13747	12701	1047	_	CDS	ND2	NADH dehydrogenase subunit 2	_	ATG	TAG
13816	13748	69	_	tRNA	trnM(cau)	tRNA-Met	CAT	_	1-
13816	13886	71	+	tRNA	trnQ(uug)	tRNA-Gln	TTG	_	_
13971	13900	72	_	tRNA	trnl(gau)	tRNA-Ile	GAU	-	_
14947	13970	978	_	CDS	ND1	NADH dehydrogenase subunit 1	_	ATG	AGG
15030	14957	74	_	tRNA	trnL(uaa)	tRNA-Leu	TAA	_	
16634	15030	1605	_	rRNA	I-rRNA	16S ribosomal RNA	_	-	_
16706	16635	72	_	tRNA	trnV(uac)	tRNA-Val	TAC	-	_
17681	16706	976	-	rRNA	s-rRNA	12S ribosomal RNA	_	_	
17750	17681	70	_	tRNA	trnF(gaa)	tRNA-Phe	GAA	_	-

Table 3. Number of short sequence repeats in mitochondrial genome of *Sarcogyps calvus*. Abbreviations: MRS, monomeric repeated sequences; DRS, dinomeric repeated sequences; TriRS, trimeric repeated sequences; TetRS, tetrameric repeated sequences.

Region	MRS	DRS	TriRS	TetRS	Microsatellite sequences	Total
(unidentified region)	7	0	4	0	2	13
ATP6	7	18	14	3	3	45
ATP8	4	4	5	1	0	14
COX1	11	56	28	7	2	104
COX2	7	15	14	4	1	41
COX3	10	27	16	2	1	56
CR1	22	46	15	3	2	88
CR2	24	44	16	2	4	90
СҮТВ	18	37	16	6	3	80
I-rRNA	43	49	18	5	0	115
ND1	17	38	15	2	3	75
ND2	18	28	20	4	5	75
ND3	3	9	3	2	0	17
ND4	27	41	34	2	1	105
ND4L	1	10	3	2	0	16
ND5	32	53	41	7	2	135
ND6	14	10	12	5	5	46
s-rRNA	20	22	10	3	1	56
trnA	1	1	0	1	0	3
trnC	0	3	0	0	0	3
trnD	1	1	0	0	0	2
trnE(uuc)	2	1	1	0	0	4
trnG	2	5	2	0	0	9
trnH(gug)	1	1	0	2	0	4
trnl	0	4	0	0	0	4
trnK	1	2	4	0	0	7
trnL	1	3	0	0	0	4
trnL(uag)	1	0	0	0	0	1
trnM	2	0	2	0	0	4
trnN	1	0	0	1	0	2
trnP(ugg)	2	2	0	0	0	4
trnQ	2	3	2	0	0	7
trnR	0	1	0	1	0	2
trnS	0	1	1	2	0	4
trnS(gcu)	2	1	0	0	0	3
trnT(ugu)	2	4	0	0	0	6
trnV	0	1	3	2	0	6
trnW	1	2	0	0	0	3
trnY	4	0	0	0	0	4
Total	311	543	299	69	35	1257

ND4 (1 site), ND4L (1 site) and ND6 (5 sites). Among these, Pro was the most frequently substituted amino acid, with 7 substitutions across ATP8, CYTB, ND1, ND3 and ND6, followed by Cys (5 sites) and Gly (4 sites) (Table 4). In the Old World vulture clade, 43 amino acid substitution sites were identified between S. calvus and other Gypini. The largest number of substitutions was found in CYTB (9 sites), while the smallest number occurred in COX1, COX2 and ND4L (1 site each) (Table 5). Unique amino acid chemical groups were found at 15 sites in S. calvus, with the largest number located in ND5 (5 sites). Pro was the most frequently substituted amino acid in this group (3 sites in ATP8, CYTB and ND4), with Gly ranking second (1 site in ND5) (Table 5).

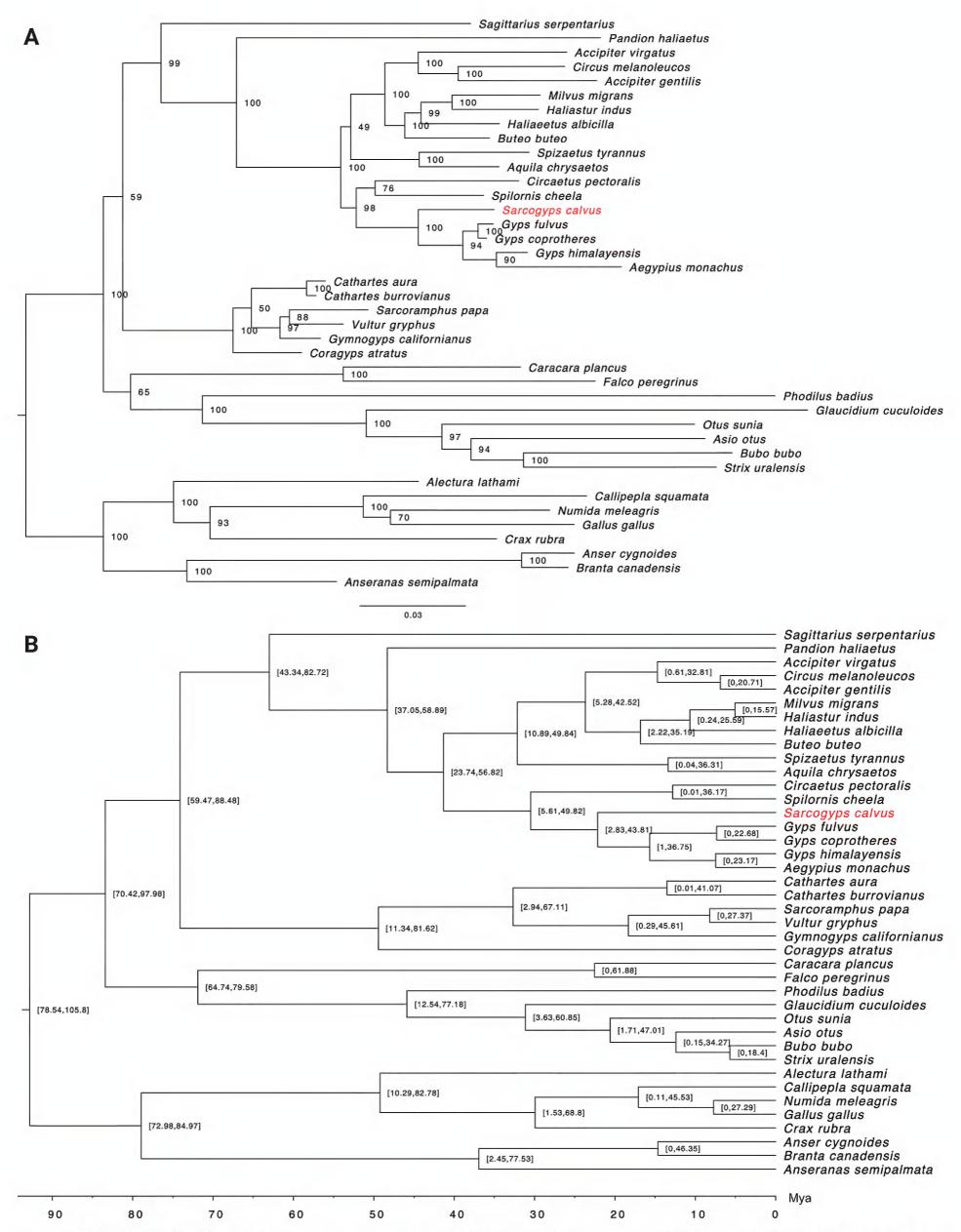


Figure 3. Maximum-likelihood phylogenetic trees based on amino acid alignments of 13 conserved protein-coding genes from the mitochondrial genomes of *Sarcogyps calvus* and 39 other bird species, with species from the orders Galliformes and Anseriformes used as outgroups **A** phylogram indicating bootstrap support values based on 1,000 replicates at each node **B** time calibrated phylogeny with the estimated minimum and maximum divergence times (millions of years ago, Mya) are indicated at each node. *Sarcogyps calvus* is in red.

Table 4. Amino acid substitution with different side chain property between Old World vultures (OWVs) and New World vultures (NWVs).

Gene	Position	OWV	NWV	Side chain property		
	_			OWV	NWV	
ATP8	7	A	N	Hydrophobic	Polar uncharged	
	30	Р	S	Special case	Polar uncharged	
COX1	1	G	Α	Special case	Hydrophobic	
	3	S	F	Polar uncharged	Hydrophobic	
COX2	4	Н	N	Positive	Polar uncharged	
	43	Α	Т	Hydrophobic	Polar uncharged	
	90	N	D	Polar uncharged	Negative	
	156	N	G	Polar uncharged	Special case	
	161	Α	S	Hydrophobic	Polar uncharged	
	166	Α	Т	Hydrophobic	Polar uncharged	
COX3	152	M	Т	Hydrophobic	Polar uncharged	
	224	N	K	Polar uncharged	Positive	
CYTB	5	Р	I	Special case	Hydrophobic	
	376	M	Т	Hydrophobic	Polar uncharged	
	378	С	Υ	Special case	Hydrophobic	
	381	Т	K	Polar uncharged	Positive	
VD1	9	Н	Υ	Positive	Hydrophobic	
	79	Т	I	Polar uncharged	Hydrophobic	
	81	M	Т	Hydrophobic	Polar uncharged	
	160	V	Т	Hydrophobic	Polar uncharged	
	171	Α	Т	Hydrophobic	Polar uncharged	
	173	Т	А	Polar uncharged	Hydrophobic	
	189	Т	А	Polar uncharged	Hydrophobic	
	260	Q	E	Polar uncharged	Negative	
	263	Т	Р	Polar uncharged	Special case	
	312	1	Т	Hydrophobic	Polar uncharged	
	323	С	Υ	Special case	Hydrophobic	
VD2	5	Т	А	Polar uncharged	Hydrophobic	
	56	T	Α	Polar uncharged	Hydrophobic	
	65	Т	А	Polar uncharged	Hydrophobic	
	140	V	Т	Hydrophobic	Polar uncharged	
	185	А	S	Hydrophobic	Polar uncharged	
	229	Т	М	Polar uncharged	Hydrophobic	
	299	Н	Υ	Positive	Hydrophobic	
ND3	7	Т	М	Polar uncharged	Hydrophobic	
	19	1	Т	Hydrophobic	Polar uncharged	
	64	L	Р	Hydrophobic	Special case	
	77	T	Р	Polar uncharged	Special case	
ND4	8	Т	M	Polar uncharged	Hydrophobic	
	40	Н	Q	Positive	Polar uncharged	
	57	C	G	Special case	Special case	
	63	S	A	Polar uncharged	Hydrophobic	
	112	A	T	Hydrophobic	Polar uncharged	
	170	Н	Q	Positive	Polar uncharged	

0	Position	owv	NWV	Side chain property		
Gene	Position			OWV	NWV	
	171	I	Т	Hydrophobic	Polar uncharged	
	195	L	S	Hydrophobic	Polar uncharged	
	201	М	Т	Hydrophobic	Polar uncharged	
ND4L	11	S	А	Polar uncharged	Hydrophobic	
	16	С	S	Special case	Polar uncharged	
	43	Т	Α	Polar uncharged	Hydrophobic	
ND5	16	А	Т	Hydrophobic	Polar uncharged	
	61	S	М	Polar uncharged	Hydrophobic	
	71	I	Т	Hydrophobic	Polar uncharged	
	332	Т	М	Polar uncharged	Hydrophobic	
	350	А	N	Hydrophobic	Polar uncharged	
	382	М	Т	Hydrophobic	Polar uncharged	
	407	А	Т	Hydrophobic	Polar uncharged	
	438	M	Т	Hydrophobic	Polar uncharged	
	597	Т	[Polar uncharged	Hydrophobic	
ND6	36	G	A	Special case	Hydrophobic	
	50	Р	S	Special case	Polar uncharged	
	53	А	S	Hydrophobic	Polar uncharged	
	73	С	S	Special case	Polar uncharged	
	78	L	Р	Hydrophobic	Special case	
	107	Е	G	Negative	Special case	
	126	V	S	Hydrophobic	Polar uncharged	
	140	W	R	Hydrophobic	Positive	

Discussion

The present study has, for the first time, characterized the complete mitogenome of S. calvus and compared it with 39 other avian mitogenomes. The mitogenome of S. calvus included 13 PCGs, 22 tRNA genes, two rRNA genes and two putative CR regions, consistent with the mitogenomes of the Gyps himalayensis and Aepygius cinereus (Li et al. 2015; Jiang et al. 2019). The total length of the PCG region of S. calvus was 11,407 base pairs (bp), which fell within the range observed in other members of Accipitriformes (11,377-11,920 bp). We observed that mitogenomes are subject to weaker translational selection compared to nuclear genomes (dos Reis et al. 2004). Regarding the GTG initiation codon of COX1, a previous study has reported the utilization of GTG as a start codon in the ribosomal protein L16 (rpl16) gene in some plant mitochondria (Bock et al. 1994). Additionally, RNA editing could also be a factor, as it has been observed in chicken mitochondria (Yokobori and Pääbo 1997). In the context of RSCU values, we identified AGG and AGA as the preferred stop codons in the mitogenome of S. calvus, with the RSCU values of 1.7 and 1.52, respectively. AGG has been identified as a stop codon for the NADH dehydrogenase 1 (ND1) and cytochrome c oxidase subunit 1(COX1) mitochondrial genes in the cinereous vulture and Himalayan griffon (Li et al. 2015; Jiang et al. 2019). Similarly, AGA has been shown to function as a stop codon for the NADH dehydrogenase subunit 3 (NADH3) and NADH dehydrogenase subunit 5 (NADH5) mitochondrial genes in the ostrich (Härlid et al. 1997). In contrast, UAA and

Table 5. Amino acid substitution with different side chain property between Old World vultures (OWVs) and New World vultures (NWVs).

Gene	Position	owv	NWV	Side chain property		
				OWV	NWV	
ATP6	31	F	I	Hydrophobic	Hydrophobic	
	83	I	I	Hydrophobic	Hydrophobic	
	139	I	I	Hydrophobic	Hydrophobic	
ATP8	24	I	l l	Hydrophobic	Hydrophobic	
	50	S	Р	Polar uncharged	Special case	
COX1	468	М	M	Hydrophobic	Hydrophobic	
COX2	70	I	I	Hydrophobic	Hydrophobic	
СҮТВ	15	I	[Hydrophobic	Hydrophobic	
	26	Р	S	Special case	Polar uncharged	
	47	L	L	Hydrophobic	Hydrophobic	
	213	1	V	Hydrophobic	Hydrophobic	
	220	Р	Р	Special case	Special case	
	307	F	F	Hydrophobic	Hydrophobic	
	310	K	К	Positive	Positive	
	321	L	L	Hydrophobic	Hydrophobic	
	370	Т	L	Polar uncharged	Hydrophobic	
ND1	15	S	S	Polar uncharged	Polar uncharged	
	179	L	L	Hydrophobic	Hydrophobic	
ND2	19	1	1	Hydrophobic	Hydrophobic	
	22	S	S	Polar uncharged	Polar uncharged	
	122	S	S	Polar uncharged	Polar uncharged	
	325	Т	Т	Polar uncharged	Polar uncharged	
	327	I	Т	Hydrophobic	Polar uncharged	
	335	I	L	Hydrophobic	Hydrophobic	
ND3	1	I	I	Hydrophobic	Hydrophobic	
	108	T	N	Hydrophobic	Polar uncharged	
ND4	43	L	L	Hydrophobic	Hydrophobic	
	90	A	T	Hydrophobic	Polar uncharged	
	183	Н	Р	Positive	Special case	
	263	T	T	Polar uncharged	Polar uncharged	
	357	T	T	Polar uncharged	Polar uncharged	
	394	i		Hydrophobic	Hydrophobic	
	418	<u>·</u> 	T	Polar uncharged	Polar uncharged	
 ND4L	73	<u>'</u> 	т	Polar uncharged	Polar uncharged	
ND5	30	 T	 	Polar uncharged	Polar uncharged	
	74		T	Hydrophobic	Polar uncharged	
	291	T	T	Polar uncharged	Polar uncharged	
	404	Y	Y	Hydrophobic	Hydrophobic	
	434	'	G	Negative	Special case	
	600		ı	Hydrophobic	Hydrophobic	
ND6	3	Λ	T			
ND6	-	A A		Hydrophobic	Polar uncharged	
	142 166	A	А	Hydrophobic Hydrophobic	Hydrophobic Hydrophobic	

UAG stop codons exhibited a negative bias, with RSCU values of 0.64 and 0.14, respectively. Additionally, we detected a negative bias against guanine at the third codon position across all 13 PCGs, consistent with findings in the cinereous vulture and Himalayan griffon (Li et al. 2015; Jiang et al. 2019).

Our results corroborate the position of *S. calvus* within the Old World vulture clade (Gypini), consistent with previous studies (Seibold and Helbig 1995; Lerner and Mindell 2005; Mindell et al. 2018; Khatri et al. 2019; Catanach et al. 2024). We estimated that S. calvus diverged from its sister clade (Gyps and Aegypius) approximately 22 Mya, while the divergence between Gypini and Cathartidae was estimated to have occurred around 74.1 Mya. This estimate closely aligns with previous findings (De Panis et al. 2021). Our analysis of amino acid substitutions, particularly those involving different chemical groups, suggests that these changes could potentially influence protein structure and function (Teng et al. 2010). Substitutions involving Cys, Pro, and Gly are particularly significant due to their unique roles in protein structure and stability. In this study, we observed transitions from Cys, which forms disulfide bonds critical for protein stability, to hydrophobic residues, potentially affecting protein folding and stability (Zavodszky et al. 2001; Alvares et al. 2013). Additionally, we detected changes involving Pro, known to restrict backbone flexibility, and Gly, which contributes to protein folding due to its small size, and may disrupt protein dynamics (Wilman et al. 2014; Senthil et al. 2019). We also noticed substitutions from hydrophobic to polar uncharged residues, such as Ser and Thr, across mitochondrial genes. These residues enhance hydrogen bonding and stability in aqueous environments, although transitions between similar residues (e.g., Ser to Thr) likely have minimal structural impact (Saeki et al. 2013). The observed amino acid changes may reflect functional adaptations and divergence within Gypini, with implications for mitochondrial function and the conservation of S. calvus. Future studies should explore these findings using structural modeling to better understand their impact.

Conclusions

Our study documents the characteristics of the complete mitogenome of *S. calvus*. Phylogenetic analysis corroborated its evolutionary relationships within Accipitriformes. *S. calvus* was most closely related to a clade formed by *Aegypius monachus* and species of *Gyps*. Additionally, we identified conserved amino acid changes between Gypini and Cathartidae, as well as unique amino acid substitutions specific to the *S. calvus*. These findings enhance our understanding of the evolutionary history and functional genomics of this critically endangered species.

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Additional information

Conflict of interest

The authors have declared that no competing interests exist.

Ethical statement

Sampling was conducted in compliance with the ethical guidelines under the Chulalongkorn University Animal Care and Use Committee (CU-ACUC), Thailand (approval number 2131005).

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Author contributions

Wannapol Buthasane: investigation; formal analysis; data curation; methodology; visualization; funding acquisition; writing – original draft. Sithichoke Tangphatsornruang: methodology; software; validation; formal analysis. Piroon Jenjaroenpun: data curation; formal analysis; investigation; methodology; software; validation. Thidathip Wongsurawat: data curation; formal analysis; investigation; methodology; software; validation. Saowaphang Sanannu; resources. Vorasuk Shotelersuk: methodology; software. Gunnaporn Suriyaphol: Conceptualization; funding acquisition; project administration; validation; supervision; writing – original draft; writing – review and editing.

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Data availability

All of the data that support the findings of this study are available in the main text or Supplementary Information.

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Supplementary material 1

Additional tables

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Explanation note: **table S1.** Pairwise genetic distances of *Sarcogyps calvus* and related species. Abbreviations: OWV, Old World vulture; NWV, New World vulture. **table S2.** Conserved amino acid substitution between Old World vultures (OWV) and New World vultures (NWV).

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